Relaxin-3 Inputs Target Hippocampal Interneurons and Deletion of Hilar Relaxin-3 Receptors in ‘Floxed-RXFP3’ Mice Impairs Spatial Memory

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ABSTRACT

Hippocampus is innervated by γ -aminobutyric acid (GABA) 'projection' neurons of the *nucleus incertus* (NI), including a population expressing the neuropeptide, relaxin-3 (RLN3). In studies aimed at gaining an understanding of the role of RLN3 signaling in hippocampus via its $G_{i/o}$ -protein-coupled receptor, RXFP3, we examined the distribution of RLN3-immunoreactive nerve fibres and RXFP3 mRNA-positive neurons in relation to hippocampal GABA neuron populations. RLN3-positive elements were detected in close-apposition with a substantial population of somatostatin (SST)- and GABA-immunoreactive neurons, and a smaller population of parvalbumin- and calretinin-immunoreactive neurons in different hippocampal areas, consistent with the relative distribution patterns of RXFP3 mRNA and these marker transcripts. In light of the functional importance of the dentate gyrus (DG) hilus in learning and memory, and our anatomical data, we examined the possible influence of RLN3/RXFP3 signaling in this region on spatial memory. Using viral-based Cre/*LoxP* recombination methods and adult mice with a 'floxed' *Rxfp3* gene, we deleted *Rxfp3* from DG hilar neurons and assessed spatial memory performance and affective behaviors. Following infusions of an AAV^(1/2)-Cre-IRES-eGFP vector, Cre expression was observed in DG hilar neurons, including SST-positive cells, and *in situ* hybridization histochemistry for RXFP3 mRNA confirmed receptor deletion relative to levels in floxed-RXFP3 mice infused with an AAV^(1/2)-eGFP (control) vector. RXFP3 depletion within the DG hilus impaired spatial reference memory in an appetitive T-maze task reflected by a reduced percentage of correct choices and increased time to meet criteria, relative to control. In a continuous spontaneous alternation Y-maze task, RXFP3-depleted mice made fewer alternations in the first minute, suggesting impairment of spatial working memory. However, RXFP3-depleted and control mice displayed similar locomotor activity, anxiety-like behavior in light/dark box and elevated-plus maze tests, and learning and long-term memory retention in the Morris water maze. These data indicate endogenous RLN3/RXFP3 signaling can modulate hippocampal-dependent spatial reference and working memory via effects on SST interneurons, and further our knowledge of hippocampal cognitive processing.

INTRODUCTION

The hippocampus plays an important role in learning and memory, in both rodents and humans (O'Keefe and Nadel, 1978; Burgess et al., 2002). The hilus (or hilar region) of the dentate gyrus (DG) contains a diverse population of interneurons, which are thought to be crucial in mediating these roles (Freund and Buzsaki, 1996). Interneurons within the DG hilus (as well as hippocampal CA1 and CA3 fields) receive GABAergic and cholinergic inputs from the medial septum/diagonal band of Broca (MS/DB) (Freund and Antal, 1988; Dutar et al., 1995; Vertes and Kocsis, 1997), which together with a number of other subcortical structures (e.g. supramammillary nucleus (Kocsis and Vertes, 1997; Ruediger et al., 2011), median raphe (MR) (Vertes et al., 1999; McKay et al., 2013) constitute the 'septohippocampal system' (Freund and Antal, 1988; Dutar et al., 1995; Vertes and Kocsis, 1997). The majority of GABAergic septal projection neurons terminate on DG hilus interneurons (Freund and Antal, 1988; Amaral et al., 2007) participating in the control of hippocampal theta oscillations (Vertes and Kocsis, 1997; Vertes, 2005; Lubenov and Siapas, 2009), and the processing of cognitive and spatial maps (O'Keefe, 1993; Leutgeb et al., 2005; Schiller et al., 2015).

In addition to these well characterized forebrain subcortical sources of afferent input, brainstem GABA projections to the hippocampus have been characterized (see Brown and McKenna, 2015, for review). GABAergic projections from the nucleus incertus (NI) (Nunez et al., 2006; Olucha-Bordonau et al., 2012; Ma et al., 2013; Sanchez-Perez et al., 2015) and nucleus of Gudden (VTg) (Bassant and Poindessous-Jazat, 2001; Kocsis et al., 2001; Bassant and Poindessous-Jazat, 2002) can regulate hippocampal theta rhythm (4-8 Hz in rodents and 4-12 Hz in humans), which is important for spatial navigation and memory formation. The NI, which sits within the dorsal tegmentum, sends strong GABA-positive projections to the MS/DB, hippocampus, supramammillary nucleus and MR nucleus, amongst other hypothalamic and limbic circuits that influence hippocampal activity (Goto et al., 2001; Olucha-Bordonau et al., 2003). Stimulation of the NI in urethane-anesthetized rats increased theta and decreased delta rhythm activity in the hippocampus, whereas electrolytic lesions of the NI abolished hippocampal theta rhythm (Nunez et al., 2006).

More recently, it was reported that lidocaine infusion into the NI impaired long-term spatial memory of adult rats in a Morris water maze (MWM) (Nategh et al., 2015) and delayed learning and impaired retention in a passive avoidance learning (PAL) task (Nategh et al., 2016), which are both hippocampal-dependent memory tasks. In the latter study, perforant path-DG short-term synaptic plasticity was also examined upon NI inactivation, both before a paired-pulse stimulation, and before or after tetanic stimulation, in freely-moving rats; revealing that NI inactivation did not change the perforant path-DG granule cell synaptic input, but decreased the excitability of DG granule cells, consistent with an effect of normal NI activity to inhibit the inhibitory interneurons in the DG and disinhibit granule cells (Nategh et al., 2016).

A large population of NI neurons express abundant levels of the neuropeptide relaxin-3 (RLN3) (Bathgate et al., 2002; Burazin et al., 2002), which acts via its cognate $G_{i/o}$ -protein coupled receptor, relaxin family peptide 3 receptor (RXFP3) (Liu et al., 2003; Bathgate et al., 2006). RLN3 neurons in the NI co-express glutamate decarboxylase (GAD), the enzyme involved in converting glutamate to GABA (Ma et al., 2007). RLN3-positive efferents overlap extensively with NI forebrain projections (Ma et al., 2007; Ma et al., 2009c; Smith et al., 2010), and in a GAD2-Cre reporter mouse, injection of a Cre-dependent viral vector expressing enhanced yellow fluorescent protein (eYFP) into the NI produced strong eYFP colocalization with RLN3-IR in NI neurons, and in hippocampus (including DG hilus) extensive eYFP-IR was observed, suggesting RLN3- and GAD-positive neurons in the NI provide projections to the hippocampus (S Ma, personal communication). RLN3-positive fibres make close contacts with MS/DB neurons which project to the hippocampus, including populations that express choline acetyltransferase (ChAT) or GAD67 and/or parvalbumin (PV) cells (Olucha-Bordonau et al., 2012), which are known to act as ‘pacemaker’ cells for hippocampal theta rhythm. Furthermore, the distribution of RLN3 fibres strongly overlaps with that of RXFP3 mRNA in the MS/DB and hippocampus, and other forebrain areas (Tanaka et al., 2005; Ma et al., 2007; Smith et al., 2010). Injection of a RXFP3-selective agonist peptide (R3/I5; Liu et al., 2005) into the MS increased hippocampal theta rhythm in urethane-anesthetized rats, which was significantly attenuated by prior injection of a selective RXFP3 antagonist (R3(BA23-27)R/I5, Kuei et al., 2007). In conscious rats, R3/I5 injection into the MS increased hippocampal theta rhythm in a home cage environment, whereas injections of R3(BA23-27)R/I5 dose-dependently reduced hippocampal theta rhythm in rats exploring a novel, enriched context, and impaired performance in a spontaneous alternation task (SAT) (Ma et al., 2009a). However, the neurochemical mechanism(s) by which the RLN3/RXFP3 system modulates hippocampal activity are largely unknown and there are no studies of this system in mice.

In the present studies designed to address this gap in knowledge, the first aim was to determine which hippocampal neurons are potentially regulated by RLN3/RXFP3 signaling. We initially examined the regional distribution of RXFP3 mRNA in the mouse hippocampus, but due to the current unavailability of a fully validated RXFP3 antibody, double-label immunohistochemistry studies to neurochemically phenotype RXFP3-positive neurons were not possible. Therefore, we conducted a series of double- and triple-label immunohistochemical experiments using a characterized monoclonal RLN3 antisera and antisera raised against key markers expressed by the hippocampus, including GABA, the neuropeptide somatostatin (SST), and the calcium-binding proteins PV and calretinin (CR). Next we assessed the behavioral role of the RLN3/RXFP3 system within the DG hilus in ‘floxed-RXFP3’ mice, in which the endogenous RXFP3 gene is flanked by two *loxP* sites (‘floxed’). In these floxed-RXFP3 mice, RXFP3 can be ‘conditionally’ deleted from neurons within discrete brain regions of adult mice by local viral delivery of the Cre recombinase restriction enzyme, which detects

loxP sites and deletes the stretch of DNA between them (Ryding et al., 2001). Thus, this approach provides temporal control and spatial specificity of receptor deletion.

These studies have revealed that RLN3-positive fibres terminate on a population of SST- and GABA-positive neurons, and a smaller population of PV- and CR- positive neurons. Appositions were observed most notably with SST-positive neurons in the DG hilus, as well as in CA1 and CA3 fields. Interestingly, conditional hippocampal RXFP3 depletion in the DG hilus impaired spatial reference memory in mice, in an appetitive T-maze reference memory task and spatial working memory in a continuous SAT in a Y-maze. We also assessed anxiety-like behavior, based on the key functional influence of the ventral hippocampus on stress and anxiety (Bannerman et al., 2004; Engelmann et al., 2006), and on the substantial number of neuroanatomical and functional studies that suggest a role for RLN3/RXFP3 signaling in responses to stress and anxiety (Banerjee et al., 2010; Smith et al., 2011; Watanabe et al., 2011; Shabanpoor et al., 2012; Ma et al., 2013; Ryan et al., 2013; Smith et al., 2014). Notably however, anxiety-like behavior was unaltered, suggesting a likely regional functional specificity of RLN3/RXFP3 signaling within the hippocampus, although the viral spread did not extend across the entire dorsoventral extent of the DG hilus in the ventral hippocampus, therefore likely reducing any potential effect on anxiety-like behavior. Taken together, these findings suggest that *endogenous* RLN3/RXFP3 signaling modulates neurons within the DG hilus, particularly a population of GABA/SST-expressing neurons, to promote spatial reference and working memory (Andrews-Zwilling et al., 2012).

MATERIALS AND METHODS

Mice

'Floxed-RXFP3' mice were generated by Gen-O-way (Paris, France), and kindly supplied by Janssen Pharmaceutical Companies of Johnson & Johnson (La Jolla, CA, USA), and bred within The Florey Institute of Neuroscience and Mental Health Animal Facility. These mice possess a 5'-*loxP* site located 1.4 kb upstream of the RXFP3 5'UTR, while a 3'-*loxP* site is located directly after the STOP codon and upstream of the 3'UTR. Floxed-RXFP3 mice were originally generated on a 129SV/B6 mixed background before being subjected to successive rounds of backcrossing onto a C57BL/6 background for 8-10 generations, via a >99% purity speed congenic approach. All studies were conducted with approval from The Florey Institute of Neuroscience and Mental Health Animal Ethics Committee and were in accordance with ethical guidelines issued by the National Health and Medical Research Council of Australia.

Behavioral experiments were performed on male homozygous RXFP3-floxed mice between 11-20 weeks of age ($n = 12 - 13$ per group). Mice were group housed (mixed treatments, ~4 mice per box) for the duration of studies. Mice were acclimatized to behavioral rooms 24 h prior to testing, and were maintained on a 12-h light-dark cycle (lights on 0700–1900) with regular chow and water available *ad libitum*, except during a food restriction regime used during T-maze testing (see below).

Detection of RXFP3 mRNA by *In Situ* Hybridization Histochemistry

Mice were deeply anesthetized with 5% isoflurane inhalation (Delvet, Seven Hills, NSW, Australia) and then administered sodium pentobarbital (100 mg/kg. 0.1 ml, i.p.) and transcardially perfused with 0.1 M phosphate buffer (PB, 2.7 mM KCl, 11.2 mM Na₂HPO₄, 1.8 mM KH₂PO₄, pH 7.4) followed by 4% paraformaldehyde (PFA) in 0.1 M PB. Mice were decapitated and their brains removed and post-fixed in 4% PFA in 0.1 M PB for 6 days. Brains were then transferred to a solution of 4% PFA and 20% sucrose in 0.1 M PB and kept at 4°C overnight. Brains were frozen on dry ice and 25 μ m sections were cut using a sliding microtome (HistoSlide 2000, Heidelberg, Germany). Levels of RXFP3 mRNA expression in the hippocampus were analyzed by *in situ* hybridization as described (Martin and Timofeeva, 2010; Lenglos et al., 2014; Lenglos et al., 2015). An RXFP3 riboprobe was generated from the 907-bp fragment of rat RLN3 receptor RXFP3 cDNA (Gene bank NM_001008310; the probe included 274–1180 bp sequence of the complete 1431 bp rat RXFP3 cDNA; forward primer: 5'-AGCGCC-GTTTACTGGGTGGTTTG-3'; reverse primer: 5'-TGGGGTTGAGGCAGCTGTTGGAGT-3'). NCBI BLAST sequencing revealed that the cDNA for rat RXFP3 is 93% homologous to mouse RXFP3.

Double- and Triple-Label Immunohistochemistry for RLN3 and GABA, CR, PV or SST

For analysis of RLN3 in nerve fibres in relation to key neurochemical markers of hippocampal neurons, a sequential double- and triple-label immunohistochemistry protocol was used. To enhance

immunoreactivity in cell bodies, 10 week-old male mice ($n = 3$) were administered colchicine (20 μg in 5 μl) into the lateral ventricle using a stereotaxic procedure (described below), at a rate of 1 μl per min. Sixteen (16) h post-surgery, mice were transcardially perfused (4% PFA in 0.1 M PB) and post-fixed in 4% PFA in 0.1 M PB for 1 h at RT, then transferred to a solution of 20% sucrose in 0.1 M PB and kept at 4°C overnight. Brains were then embedded in O.C.T. (Tissue-Tek; Sakura Finetek, USA) and frozen over dry ice and stored at -80°C. Forty (40) μm coronal sections were cut and collected into 4 series spanning from -1.22 mm to -3.80 mm from Bregma using a cryostat (Cryocut 1800, Leica Microsystems, Heerbrugg, Switzerland) at -18°C and stored in cryoprotectant solution (30 % ethylene glycol, 30 % glycerol, 0.05M PB) at -20°C.

All sections were washed 3×5 min with 0.1% Triton X-100 in 0.1 M PB, and then blocked in 10% normal horse serum (NHS) in 0.1% Triton X-100, 0.1 M PB for 1 h at RT. Sections were incubated with a mouse monoclonal RLN3 antibody (HK4-144-10; Kizawa et al., 2003; Tanaka et al., 2005; Ma et al., 2013) in 2% NHS and 0.1% Triton X-100 in 0.1M PB overnight at RT (for dilutions and information about the primary antisera used, see **Table S1**). On the following day, sections were washed 3×5 min in 0.1 M PB, followed by incubation in donkey anti-mouse Alexa-594 (JIR, 715585151, 1:500) or donkey anti-mouse Alexa-647 (JIR, 715605151, 1:500) in 0.1 M PB for 1 h at RT. Following application of the secondary antibody for visualization of RLN3-IR, sections were washed 3×5 min with 0.1 M PB and incubated with either: (1) rabbit anti-GABA (Sigma, A2052, 1:3,000) (2) rabbit anti-PV (Abcam, ab11427, 1:1,000) and rat anti-SST (Millipore, MAB354, 1:200) (3) and rabbit anti-CR (Swant, CR7697, 1:1,000) in 2% NHS, 0.1% Triton X-100, 0.1 M PB overnight at RT. Sections were washed 3×5 min in 0.1 M PB before incubation in the following secondary antibodies (1) donkey anti-rabbit Alexa-488 (Life Tech, A21206, 1:500) (2) donkey anti-rat Alexa-488 (Life Tech, A21208, 1:500) and donkey anti-rabbit Alexa-594 (JIR, 711585152, 1:500) and (3) donkey anti-rabbit Alexa-488 (Life Tech, A21206, 1:500) in 0.1 M PB for 1 h at RT. Following the secondary antisera incubation, sections were washed 3×5 min with 0.1 M PB and mounted on glass slides using Fluoromount-G (Southern Biotech, Birmingham, AL, USA).

Resultant staining was observed under a LSM 780 Zeiss Axio Imager 2 confocal laser scanning microscope (Carl Zeiss AG, Jena, Germany). Each fluorescence channel was imaged sequentially using a 20 $\times$ objective for mosaic z stacks (1 μm intervals) and the total proportions of GABA-, SST-, PV- and CR- positive neurons contacted by RLN3-IR fibres were manually counted from the dorsal hippocampus of a representative unilateral coronal section from 3 mice (-2.54 mm from Bregma) based on the strong RLN3 fibre innervation observed at this coronal level. Mosaic z-stack images covering the entire hippocampus from each representative coronal section were obtained with 10% overlap between each image on the X-Y plane, and a defined region was set up for the z plane to scan from the bottom to the top of each image comprising the mosaic. Z-stack images (63 $\times$ objective, 0.1 μm intervals) were taken to further evaluate putative contacts between RLN3-IR nerve fibres and

hippocampal cellular markers. A putative contact was scored when an RLN3-IR fibre/terminal bouton was observed immediately adjacent to the cell body of a neuron expressing a hippocampal marker.

Adeno-Associated Viral (AAV) Vectors

The AAV mosaic serotype 1/2 with capsid from AAV1 and ITR (internal terminal repeats) AAV2 was purified and harvested as described (Ganella et al., 2013). The AAV^(1/2)-Cre-IRES-eGFP viral vector encodes Cre recombinase and an enhanced green fluorescent protein marker (eGFP) linked by an IRES sequence (internal ribosome entry site), allowing for the expression of Cre recombinase and eGFP from a single vector. The control AAV^(1/2)-eGFP construct was also packaged in a 1/2 mosaic capsid, encoding the expression of eGFP marker protein. Transcription of both constructs was driven by the chicken β -actin (sCAG) promoter. The titres were assessed visually in HEK293T cells under a fluorescent microscope and by measuring genomic copies (gc) per ml (gc/ml), using quantitative polymerase chain reaction (qPCR). The resulting titre of AAV^(1/2)-Cre-IRES-eGFP and the control AAV^(1/2)-eGFP vector were approximately 2×10^{11} gc/ml.

Stereotaxic Surgery

Mice were initially anesthetized with 5% isoflurane inhalation (Delvet) mixed with oxygen till loss of righting reflex, then secured in a small animal stereotaxic frame (Kopf Instruments, Tujunga, CA, USA) with 1.5-2% isoflurane delivered through a small animal nose cone. Analgesic (Meloxicam 20mg/kg; Troy Laboratories, Smithfield, NSW, Australia) was then administered intraperitoneally (i.p., 0.1 ml), and eyes were moistened with lubricating eye ointment (Lacri-Lube, Allergen, New Jersey, USA). A small midline incision was made to expose the skull, which was then cleaned/dried with 6% hydrogen peroxide. AAV^(1/2)-Cre-IRES-eGFP viral vector or control AAV^(1/2)-eGFP viral vector was loaded into a glass capillary injector which was connected to polyethylene tubing and a 10 μ l Hamilton syringe (0.46 mm diameter, Harvard Apparatus, Holliston, MA, USA), and mounted on an infusion pump (PicoPlus, Harvard). Small holes were drilled through the skull above the injection sites, and the glass capillary was lowered to the following coordinates from Bregma based on the mouse brain atlas (Paxinos and Franklin, 2001): anterior-posterior, -2.3 mm; medial-lateral, -1.60 mm; dorsal-ventral, -2.1 mm; adjusted proportionally for skull size, determined by bregma-interaural distance). Viral vectors were infused at a rate of 0.1 μ l/min to a total volume of 1 μ l. After each infusion, the injector was left in place for 7 min, retracted 1.0 mm, and left for another min to minimize deposition of virus in the injection tract, before being slowly removed. After surgery, the skin incision site and holes in the skull were closed/sealed with glue (super glue, Daiso, Hiroshima, Japan) and a single suture. After surgery mice were placed in recovery chambers (30°C, Thermacage, Datasand Ltd., Manchester, UK) for ~1 h, and left to recover for 3 weeks to allow for Cre expression, RXFP3 gene deletion, and subsequent RXFP3 protein deletion.

Validation of Cre Expression (Cre-IR) Within SST-Positive Neurons

In studies to assess Cre-recombinase expression in the targeted area and the expression of Cre within SST-positive neurons, coronal sections (40 μ m) were washed 3×5 min with 0.1% Triton X-100 in 0.1 M PB, and then blocked in 10% NHS in 0.1% Triton X-100 and 0.1 M PB for 1 h at RT. Tissue were then incubated with rabbit polyclonal Cre antibody (Novagen, 69050-3, 1:500, $n = 3$) and rat monoclonal anti-SST (Millipore, MAB354, 1:200, $n = 3$) in 2% NHS and 0.1% Triton X-100 overnight at RT. On the following day, sections were washed 3×5 min in 0.1 M PB, followed by incubation in anti-rabbit Alexa-594 secondary antibody (Jackson ImmunoResearch Laboratories, Address, USA, 120330, 1:500) and anti-rat Alexa-594 secondary antibody (1:500) in 0.1 M PB. Sections were then washed 3×5 min with 0.1 M PB and mounted on glass slides using Fluoromount-G (Southern Biotech, Birmingham, AL, USA). The resulting staining was observed under a Zeiss Axio Imager 2 confocal laser-scanning microscope (Carl Zeiss AG, Jena, Germany).

Validation of RXFP3 mRNA Deletion

RXFP3 mRNA loss following AAV^(1/2)-Cre-IRES-eGFP injection, was measured in mice injected bilaterally with AAV^(1/2)-Cre-IRES-eGFP ($n = 5$), or with AAV^(1/2)-eGFP ($n = 3$). Twenty-one days after stereotaxic surgery, mice were transcardially perfused (4% PFA in 0.1 M PB). For quantitative analyses, slides were examined with dark-field microscopy using an Olympus BX61 microscope (Olympus Canada, Richmond Hill, ON, Canada). Images were acquired with a DVC-2000C digital camera (Thorslabs Scientific Imaging, Austin, TX, USA) and analyzed with Stereo Investigator software (MBF Bioscience, Williston, VT, USA). Brain sections from bregma -1.7 mm to -2.7 mm (based on a stereotaxic atlas of mouse brain, (Paxinos and Franklin, 2001)) were used for quantitative analyses. The optical density (OD) was obtained by defining the contour of the region of interest (DG and CA3), and three background contours (regions without positive hybridization signal) were collected to correct for the average background signal. This analysis was taken from 3-5 sections on each side of the brain for bilaterally injected mice, and from 7-12 sections from unilaterally injected mice. The mean OD of AAV^(1/2)-Cre injected tissue was normalized to the mean value of RXFP3 mRNA in AAV^(1/2)-GFP injected tissue, to obtain the relative level of RXFP3 mRNA for each section.

Validation of Cre Targeting of Dentate Gyrus Hilus

Following confirmation that local hippocampal injection of AAV^(1/2)-Cre-IRES-eGFP viral vector resulted in Cre expression and deletion of RXFP3 mRNA in floxed-RXFP3 mice, in subsequent studies, the presence of Cre-IR throughout the DG hilus was taken as sufficient evidence of adequate RXFP3 deletion within this structure. For all histological analyses a 1-in-3 series of brain sections was collected spanning from -1.22 mm to -3.80 mm from Bregma (Paxinos and Franklin, 2001), and only mice which had consistent expression of Cre-IR throughout the DG hilus of dorsal hippocampus were included in behavioral analyses. Of the 16 mice that received bilateral AAV^(1/2)-Cre-IRES-eGFP

injections, 4 mice were excluded from the analyses due to a lack of transduced neurons present in the DG hilus. No viral spread was observed outside the DG and CA3.

Behavioral Tests

All behavioral experiments were conducted during the light phase, between 0900-1700 h, and were conducted from 21 days after surgery in the following order: (1) Automated locomotor cell; (2) Morris water maze (MWM); (3) Light/dark box; (4) Elevated plus maze; (5) Continuous spontaneous alternation test, and; (6) Appetitive spatial reference T-maze test. All mice were sacrificed 3 days after behavioral testing for histological analyses.

Automated Locomotor Cell

Mice were tested in a 27 × 27 cm automated locomotor cell (Med Associates, Fairfax, Vermont, USA), illuminated by 70 lux light, for a total duration of 1 h. The total distance travelled was tracked by a photobeam array. Data was analyzed using Activity Monitor, v.9.02 software (v.6.02; Med Associates).

Morris Water Maze (MWM)

In a test of hippocampus-dependent spatial reference long-term memory, mice were trained in a circular pool (1.2 m diameter) to locate a hidden platform (10 cm diameter) which was submerged ~0.5-1 cm beneath opaque water, made using non-toxic white paint and maintained at 18-22°C. The pool was surrounded by distal extra-maze cues. The acquisition phase consisted of 6 consecutive training days with 4 trials per day, with an inter-trial interval (ITI) of 30 min. For each trial, mice were placed in from a semi-random starting location (N, NW, E, SE) facing the pool wall, and were allowed to swim until they found the hidden platform or for a maximum of 30 s. If a mouse failed to locate the platform within the allocated time, the experimenter guided the mouse to the platform. The mouse then remained on the platform for 30 s, before being returned to its home cage, and allowed to dry under a heat lamp. For the acquisition sessions, a mean latency to locate the target quadrant was calculated for each mouse by averaging the latency to reach the platform across all 4 trials. Twenty-four (24) h after the last training day, the platform was removed from the target quadrant and the mice were allowed to swim freely for a duration of 120 s. The percent time spent in the target quadrant and the number of platform crossings was calculated.

Anxiety-like Behavior - Light/Dark Box

Mice were placed in the automated cells described above with half of the locomotor cell fitted with a black Perspex box, which was impermeable to visible light. The other half was exposed to a light-emitting diode array, illuminated the light side to ~700 lux in the centre and 650 lux in the corners, creating an aversive environment. For testing, mice were placed in the dark side of the apparatus for 10 min. A small opening in the Perspex box allowed access between both sides of the

cell. Time spent in and number of entries into the light side was recorded using Activity Monitor software (v.6.02; Med Associates).

Elevated Plus Maze (EPM)

The apparatus consisted of four 30 cm long and 6 cm wide arms extending from a central square ($6 \times 9 \times 6$ cm), elevated 39 cm above the ground. High (15 cm) walls enclosed two opposing 'closed' arms, while the two opposite arms were 'open'. Mice were placed in the centre of the apparatus, and were allowed to roam freely for 10 min. Movement was tracked from above using Top Scan Lite 2.0 software, and the time spent in the open arms, and the number of entries into the closed and open arms, were assessed.

Spatial Working Memory: Continuous Spontaneous Alternation Test (SAT)

The SAT was conducted using an enclosed Y-maze, with each arm 10 cm wide and 30 cm long, with 17 cm high walls. The maze was cleaned with water between tests and for testing, mice were placed in the centre of the arena facing the 'home' arm. Mice are allowed free access to the three arms of the maze for a total of 10 min during which the sequence of arm entries was recorded using Ethovision XT software. From these sequences, proportions of arms entered that were different from those previously entered were calculated to arrive at a percentage alternation score (PAS). An 'alternation' is determined from successive entries of three arms in which three different arms are entered. PAS was calculated by dividing the number of observed alternations by the number of maximum possible alternations (total number of entries -2) and multiplying the quotient by 100.

Spatial Reference Memory: Appetitive T-maze Test

Acquisition and reversal learning in a reward-based T-maze (each arm was 7.6 cm wide and each opposing arm was 26.7 cm long, with a 28.9 cm home arm, with 13 cm high walls) was tested as described (Moy et al., 2007). Acquisition of this task demands the use of allocentric spatial cues and is affected by hippocampal lesions (Deacon et al., 2002; Reisel et al., 2002; von Engelhardt et al., 2008). The maze was surrounded by distal 3-D spatial cues placed 40 cm above and outside each arm. Before testing, mice were food deprived to 85-90% of their free-feeding body weight 5 days before testing and throughout the duration of the experiment. Mice were habituated (with 4 trials per day) to the T-maze and trained to obtain a food reward (0.2 g of chocolate-flavored rice puffs) from cups located at the ends of the two choice arms. Only when the mice could readily run from the start arm to a choice arm and consume their reward was acquisition commenced. After 5 days of habituation, 10 successive trials per day were initiated. One choice arm was designated the 'reward' arm with a food reward (0.2 g chocolate-flavored rice puffs) placed at the end of the arm. The reward arm was on the left for half the mice, and on the right for the remainder, and this was randomized between treatment groups. At the beginning of each session, the mouse was placed in a start box at the bottom of the home arm. The start box door was opened, and the mouse was given the choice between entering either arm.

Following a correct choice the mouse was given time to consume the food reward and then guided back into the start box to begin the next session. If the mouse made an incorrect choice it was left in the maze for 15 s before being guided back to the start box. To meet criterion, mice were required to make 80% correct choices for 3 consecutive days before reversal training commenced, in which the reward arm was switched to the opposite arm and 10 trials per day were initiated. Testing was stopped after criterion for individual mice was met during reversal. Time spent in junction before choice were recorded using Ethovision XT software; and the number of correct choices, and number of days to meet criterion were recorded by an observer.

Statistical Analysis

All graphs and statistical analysis were conducted using Graph Pad Prism (v.6) (GraphPad, La Jolla, CA, USA). All data are expressed as mean $\pm$ SEM. Shapiro-Wilks test were employed to assess the shape of data distributions. For data sets which met the assumptions of normality, comparison of data was performed using either one-way ANOVA or two-way RM ANOVA, followed by appropriate post-hoc comparisons, as described or unpaired t-test. For data sets which did not meet the assumption of normality, the non-parametric equivalent of a t-test, the Mann-Whitney U test was used. Results were considered statistically significant if $p < 0.05$.

RESULTS

Distribution of RXFP3 mRNA and RLN3-Positive Innervation in Hippocampus

In agreement with previous studies in mouse (Smith et al., 2010) and rat (Ma et al., 2009c), the present study identified RXFP3 mRNA expression in mouse hippocampal formation (**Fig. 1A**) particularly in the DG hilus (**Fig. 1B**), across the stratum oriens layer of CA1, with less expression in the pyramidal layer of CA1 (**Fig. 1C**). RXFP3 mRNA was also detected in cells across the pyramidal, stratum oriens and radiatum layers of CA3 (**Fig. 1D**). Furthermore, the RXFP3 mRNA expression pattern overlapped the distribution of neuronal fibres and terminals containing RLN3-IR in the hippocampus (**Fig. 2A**), with abundant fibres observed in the DG hilus (**Fig. 2B**), and similar to RXFP3 mRNA levels, a smaller density of neuronal fibres containing RLN3-IR was observed in the pyramidal layer of CA1 and the stratum oriens of CA1 (**Fig. 2C**). Dense RLN3-positive fibres were detected across the pyramidal layer and radiatum of CA3 (**Fig. 2D**).

These results are also in agreement with the distribution of RXFP3 mRNA reported in the Allen Brain Institute atlas of gene expression (**Fig. 3A**). Notably, the distribution of RXFP3 mRNA and RLN3-positive nerve fibres is similar to that of a proportion or population of γ -aminobutyric acid (GABA) neurons in the hippocampus, as reflected by the pattern of GAD67 mRNA expression (**Fig. 3B**), as well as other populations of GABA neurons that express the neuropeptide, SST (**Fig. 3C**) and the calcium-binding proteins, PV (**Fig. 3D**) and CR (**Fig. 3E**), which are co-expressed with GABA in the hippocampus (Freund and Buzsaki, 1996; Jinno and Kosaka, 2002b; Baraban and Tallent, 2004).

RLN3 Innervation of Hippocampal Interneurons

For the analysis of RLN3-IR in neuronal fibres in relation to other markers of hippocampal neurons, a sequential double and triple label immunohistochemistry protocol was used with RLN3 and GABA, SST and the calcium-binding proteins, PV and CR (**Fig. 4**). Boutons containing RLN3-IR were observed in apposition to neurons positive for GABA (**Fig. 4A**), SST, PV (**Fig. 4B**; **Fig. S1**) and CR (**Fig. 4C**); and the number and proportions of the GABA-, SST-, PV- and CR- positive neurons contacted by fibres containing RLN3-IR were counted in the unilateral dorsal hippocampus of a representative coronal section (-2.54 mm from Bregma), from 3 mice (see Materials and Methods for details), based on the strong RLN3 innervation observed at this level (**Table S2**).

A high density of SST neurons was detected within the DG hilus and stratum radiatum of CA1 and CA3 with the highest density in the oriens layer of both subfields, and in the material assessed, ~26% of SST-positive neurons (6 ± 1 of 23 ± 3 neurons counted, $n = 3$) in the DG hilus received contacts from fibres containing RLN3-IR. Similarly, ~14% of SST-positive neurons (5 ± 1 of 34 ± 6 , $n = 3$) in the stratum oriens layer of CA1 and ~33% of SST-positive neurons (3 ± 2 of 8 ± 3 , $n = 3$) in the stratum radiatum of CA3 were contacted by fibres containing RLN3-IR. Furthermore, we observed RLN3-positive bouton contacts on a subpopulation of GABA-positive neurons, representing ~12% of

GABA-positive neurons (5 ± 0 of 46 ± 11 , $n = 3$) in the DG hilus and ~15% of GABA-positive neurons (6 ± 2 of 38 ± 9 , $n = 3$) in the stratum oriens layer of CA1. Notably, the number of neurons innervated by fibres containing RLN3-IR is similar for both SST and GABA populations with a total of 40 SST-positive neurons and 44 GABA-positive neurons (**Table S2**).

In contrast, neuronal fibres containing RLN3-IR were observed to innervate a smaller population of PV- and CR-positive neurons in the hippocampus, with ~23% of PV-positive neurons (1 ± 0 of 4 ± 1 , $n = 3$) in the DG hilus innervated, and ~7% of PV-positive neurons (3 ± 1 of 45 ± 8) in the oriens layer of the CA3, and ~7% of PV-positive neurons (a total of 17 of 268) in all subfields of the hippocampus. Similarly, ~9% of CR-positive neurons (2 ± 1 of 23 ± 4 , $n = 3$) in the DG hilus were innervated by fibres containing RLN3-IR, with only ~5% of CR-positive neurons (a total of 18 of 399, $n = 3$) innervated in all subfields of the hippocampus, in the material examined (**Table S2**).

***In Vivo* Validation of AAV^(1/2)-Cre-IRES-eGFP and Cre Expression in SST-positive Neurons**

Based on the reported role of SST-positive neurons in the DG hilus in learning and memory (Baratta et al., 2002; Andrews-Zwilling et al., 2010; Spiegel et al., 2013), and the appositions observed between RLN3-positive afferents and SST-positive neurons in the DG hilus, we examined the functional role of RXFP3 in this region by conditionally deleting RXFP3 in the DG hilus in adult floxed-RXFP3 mice using an AAV^(1/2)-Cre-IRES-eGFP viral vector. To validate that the AAV^(1/2)-Cre-IRES-eGFP viral vector effectively drove Cre recombinase expression *in vivo* and transduced hippocampal DG hilus SST-positive neurons, sections from adult floxed-RXFP3 mice (injected bilaterally in the DG hilus with AAV^(1/2)-Cre-IRES-eGFP) were processed for Cre- and SST-IR. Examination of these sections revealed abundant eGFP (**Fig. 5A**) and Cre-IR (**Fig. 5B**) in neurons in the DG hilus. Importantly, Cre-IR was observed in the majority of SST-positive neurons present in the DG hilus of injected mice (**Fig. 5C, D; Fig. S2**).

Confirmation of Conditional RXFP3 Deletion within the DG Hilus

Six (6) weeks after bilateral injections of AAV^(1/2)-Cre-IRES-eGFP into the DG hilus, a significant reduction in RXFP3 mRNA was observed in the DG hilus, relative to control AAV^(1/2)-eGFP bilaterally injected mice ($t_{(14)} = 3.64$, $p < 0.01$, $n = 3 - 5$ mice per group, **Fig. 6A-C**). Although Cre-IR was observed in CA3, a significant reduction in RXFP3 mRNA was not observed in this area in AAV^(1/2)-Cre-IRES-eGFP bilaterally injected mice, relative to control, AAV^(1/2)-eGFP injected mice ($t_{(14)} = 2.06$, $p = 0.058$, **Fig. 6D**). Therefore, while Cre was present in some CA3 neurons, it is presumed that an insufficient number of RXFP3 mRNA-positive neurons were transduced to produce a detectable decrease in mRNA levels. **Viral spread was observed along the rostrocaudal extent of the dorsal DG hilus and CA3 in AAV^(1/2)-Cre-IRES-eGFP treated mice, but did not reach the ventral DG hilus/CA3 regions (Fig. 6E).**

RXFP3 Deletion from DG Hilus Impaired Spatial Reference Memory in an Appetitive T-maze Task

In an appetitive T-maze task, AAV^(1/2)-Cre-IRES-eGFP-treated mice made significantly fewer correct choices compared to AAV^(1/2)-eGFP-treated control mice, but a significant increase in performance over days tested was observed in both groups (RM 2-way ANOVA, main effect of treatment, $F_{(1,21)} = 9.18$, $p = 0.0064$; main effect of day, $F_{(4,84)} = 32.6$, $p < 0.0001$; treatment $\times$ day interaction, $F_{(4,84)} = 1.89$, $p = 0.120$; Bonferroni post hoc analysis between treatments within days 1-3, $p < 0.05$, **Fig. 7A**). Although mice from both treatment groups met the criterion of 80% correct choices for 3 consecutive days, AAV^(1/2)-Cre-IRES-eGFP mice took significantly longer to meet the criterion than AAV^(1/2)-eGFP mice ($U = 35$, $p = 0.0210$, **Fig. 7B**).

When the criterion of correct choices was met during acquisition, the reward arm was swapped to the opposite arm, and reversal learning was commenced. This was done to test whether AAV^(1/2)-Cre-IRES-eGFP-treated mice displayed a resistance to alter/change a learned behavior (i.e., altered behavioral flexibility). Two-way repeated measures ANOVA did not reveal a main effect of treatment ($F_{(1,20)} = 1.79$, $p = 0.196$), nor an interaction effect ($F_{(4,80)} = 0.403$, $p = 0.806$) in % correct choice during reversal, however a main effect of day was observed ($F_{(1, 24)} = 4.46$, $p = 0.045$, **Fig. 7A**). Interestingly, during reversal, AAV^(1/2)-Cre-IRES-eGFP mice spent significantly more time in the T-maze centre junction before making a choice. No differences were observed between groups in time taken to meet the learning response criterion ($U = 66.5$, $p = 0.780$, **Fig. 7B**).

RXFP3 Deletion from DG Hilus Impaired Spatial Working Memory in a Continuous SAT in a Y-maze

In studies to measure spatial working memory, mice were assessed in a continuous SAT in a Y-maze. RXFP3 deletion in the DG hilus significantly impaired spontaneous alternation in the percent alternation score (PAS) in the first 1-min time bin ($t_{(22)} = 3.92$, $p = 0.0007$, **Fig. 8**), with a PAS score of 67.6% for AAV^(1/2)-eGFP-treated controls, and 35.8% for AAV^(1/2)-Cre-IRES-eGFP-treated mice. A t-test revealed AAV^(1/2)-eGFP mice performed significantly better than chance (PAS 50%) ($t_{(24)} = 3.02$, $p = 0.0059$), relative to AAV^(1/2)-Cre-IRES-eGFP mice that performed significantly below chance ($t_{(24)} = 2.26$, $p = 0.0329$) in the first min of the SAT test. However, over a total test time of 3 min, a significant difference between treatments was not observed ($t_{(22)} = 1.18$, $p = 0.0822$), with control mice performance only just above chance (PAS, 52.1 ± 3.35 ; $t_{(24)} = 0.630$, $p = 0.536$), and poorer performance observed for AAV^(1/2)-Cre-IRES-eGFP mice (PAS, 44.8 ± 3.54 ; $t_{(25)} = 1.80$, $p = 0.0844$).

RXFP3 Deletion from DG Hilus Did Not Alter Learning and Long-term Memory Retention in a Morris Water Maze

We next determined whether deletion of RXFP3 from the DG hilus impaired spatial learning and long-term retrieval in a MWM. A 2-way repeated measures ANOVA revealed a significant main

effect of day ($F_{(5,110)} = 35.4$, $p < 0.0001$), but no main effect of treatment was observed ($F_{(1,22)} = 0.0240$, $p = 0.878$), nor an interaction ($F_{(5,110)} = 0.272$, $p = 0.928$, **Fig. 9A**) in the latency (s) to locate the hidden platform during acquisition. Twenty-four (24) h after the last acquisition day, in the probe trial, both treatment groups spent significantly more time in the target quadrant relative to the other quadrants in the first 1 min time bin (one-way ANOVA, $F_{(7,84)} = 11.3$, $p < 0.0001$) and in the total duration of 2 min (one-way ANOVA, $F_{(7,84)} = 29.2$, $p < 0.0001$, **Fig. 9B**). Bonferroni's planned comparison tests confirmed that both AAV^(1/2)-Cre-IRES-eGFP-treated mice and AAV^(1/2)-eGFP-treated control mice spent significantly more percent time in the target quadrant than in the adjacent quadrant to the right (AAV^(1/2)-eGFP; $p < 0.0001$, AAV^(1/2)-Cre-IRES-eGFP; $p < 0.05$), the adjacent quadrant to the left ($p < 0.0001$) and to the opposite quadrant ($p < 0.0001$). No significant differences were observed between treatment groups in the percent time spent in the target quadrant in the first min ($t_{(21)} = 0.0605$, $p = 0.952$) and the total 2 min test duration ($t_{(21)} = 1.009$, $p = 0.325$). The time each group spent in the target quadrant on probe day was compared with a 'chance' value of 30 s (i.e., the amount of time a mouse would be expected to spend in the target quadrant over a total duration of 2 min if no spatial memory had been formed). Both groups spent significantly more than 25% (15 s) of their time in the target quadrant in the first min (AAV^(1/2)-GFP, $t_{(24)} = 3.91$, $p = 0.0007$; AAV^(1/2)-Cre, $t_{(24)} = 4.01$, $p = 0.0005$) and significantly more than 25% (30 s) of their time in the target quadrant in the total 2 min duration (AAV^(1/2)-GFP, $t_{(24)} = 5.80$, $p < 0.0001$; AAV^(1/2)-Cre, $t_{(24)} = 4.00$, $p = 0.0005$), suggesting intact spatial long-term memory on probe day.

RXFP3 Deletion from DG Hilus Did Not Alter General Locomotor Activity or Innate Anxiety in an EPM and L/D box

No differential effects were observed between treatments in the total distance travelled in a locomotor cell 21 days after viral injections ($t_{(23)} = 0.551$, $p = 0.587$; **Table S3**). This result was consistent across 5-min bins within a total duration of 1 h, and both groups significantly decreased their distance travelled over time (RM 2-way ANOVA, main effect of treatment ($F_{(1,23)} = 0.303$, $p = 0.587$); time ($F_{(11,253)} = 15.6$, $p < 0.0001$); treatment $\times$ time interaction ($F_{(11,253)} = 0.520$, $p = 0.889$).

Twenty-four (24) h after the probe day in the MWM, the different groups of mice were tested in an EPM to measure stress-induced anxiety-like behavior. RXFP3 deletion from DG hilus did not induce any changes relative to AAV^(1/2)-GFP control mice in the time spent in the open arms ($t_{(23)} = 0.886$, $p = 0.139$; **Table S3**), or the number of entries into the open arms ($t_{(23)} = 0.498$, $p = 0.623$). Similarly, in the L/D box, no differences were observed between groups in the total time spent in the light side ($t_{(22)} = 0.886$, $p = 0.385$; **Table S3**) or entries into the light side ($t_{(22)} = 1.528$, $p = 0.141$).

DISCUSSION

In the present study, in light of the distinct topography of RXFP3 mRNA in the mouse hippocampus, we conducted a histochemical analysis of the possible association of RLN3-positive nerve fibres with some established hippocampal neuron types; and a behavioral analysis of mice after deletion of RXFP3 from the DG hilus. The main findings were firstly, that RXFP3 mRNA was confirmed to be expressed in distinct layers of the hippocampus, with strong expression in neurons within the DG hilus, and non-principal neurons located within the pyramidal and stratum oriens layer of CA1 and across the pyramidal layer, stratum oriens and radiatum of CA3. Interestingly, neuronal fibres and terminals containing RLN3-IR were identified making consistent putative close contacts with GABA- and SST- positive neurons and, to a lesser degree, contacts with PV- and CR-positive neurons. Secondly, in this first study to manipulate RLN3/RXFP3 signaling within the hippocampus by deletion of RXFP3 from the DG hilus, we observed a strong and consistent deficiency in an appetitive reference memory task in a T-maze, and impairment in spatial working memory in a Y-maze SAT test. In contrast, deletion of RXFP3 from the DG hilus did not affect the ability of mice to learn and acquire the location of a hidden platform in a MWM test of long-term memory retention. Furthermore, hilar RXFP3 deletion did not alter anxiety-like behavior in an EPM or L/D box. Overall, this study identified key behavioral consequences of RXFP3 depletion from DG hilus and the neurochemical phenotype of the RLN3/RXFP3 targeted neurons in the hippocampus, providing new insights into the complex role of this modulatory neuropeptide system in learning and memory.

Our first goal was to assess the neurochemical anatomy of the RLN3/RXFP3 system within the hippocampus. A detailed mapping of the distribution of RXFP3 mRNA/binding sites and RLN3-IR has been completed in mouse (Smith et al., 2010), rat (Ma et al., 2009c; Olucha-Bordonau et al., 2012) and non-human primate (Ma et al., 2009c) brain and the results are consistent with the present analysis in the mouse hippocampus. Based on the overlap between the quite restricted topographic distribution of neurons expressing RXFP3 mRNA with the more extensive distribution of GABAergic interneurons, we hypothesized that RXFP3 is expressed by a subpopulation of GABA neurons in the hippocampus. This in turn predicted that neuronal fibres expressing RLN3-IR would make close contacts with GABA-positive neurons in the hippocampus, particularly in the DG hilus and the oriens layer of CA1, and this was demonstrated experimentally. Notably, RLN3-positive fibres were also found to make a significant number of contacts with SST-positive neurons in these areas. As the majority of SST neurons (~90%) in the DG hilus co-express GABA (Kosaka et al., 1988), our results indicate there is a subpopulation of RXFP3-positive neurons which co-express both GABA and SST in the hippocampus. Interestingly, RLN3-positive fibres and terminals were also observed to make close contacts with PV- and CR-positive neurons in the hippocampus, but their incidence was approximately half that for contacts with GABA- and SST-positive neurons. PV and CR label distinct, non-overlapping subsets of inhibitory hippocampal interneurons (Xu et al., 2010), therefore, it is likely that

a majority of RFXFP3-positive neurons in the hippocampus express GABA and SST, while non-overlapping smaller subpopulations of RFXFP3-positive neurons co-express PV or CR.

SST neurons constitute one of the largest groups of GABA neurons within the DG, with all these SST-positive neurons located within the hilus (Scharfman and Myers, 2012). Retrograde tracing studies in mice indicate that a large proportion (44%) of SST-positive neurons in the DG hilus project to the septal region (Zappone and Sloviter, 2001; Jinno and Kosaka, 2002a), and directly innervate GABAergic and cholinergic neurons in that region (Toth et al., 1993; Gulyas et al., 2003). In turn, SST hippocampal-septal neurons are reciprocally innervated by cholinergic and GABAergic septal neurons (Freund and Antal, 1988; Freund and Buzsaki, 1996). The high percentage of SST DG hilus neurons that innervate the septum and the high incidence of RLN3 inputs to SST- and GABA-positive neurons in the DG hilus, suggest RLN3/RFXFP3 signaling plays an important role in the hippocampo-septal and septo-hippocampal circuitry, modulating cognition and learning and memory. Therefore, it will be of interest to further investigate the precise anatomical connectivity of those neurons and circuits influenced by the RLN3/RFXFP3 system and its resultant functional impact.

However, there is a considerable amount of existing evidence regarding the functional role of SST-containing DG hilus interneurons that might inform the possible effects of RFXFP3 signaling on hippocampal activity. Activation of SST DG hilus interneurons reduces long-term potentiation (LTP) in the mouse DG (Baratta et al., 2002), a cellular process involved in learning and memory; and an age-dependent reduction in SST and GAD67 DG hilus neurons in apolipoprotein E4 (apoE4) knock-in mice correlated with learning and memory deficits in the MWM test (Andrews-Zwilling et al., 2010). Notably, a similar result was observed in a study of rat DG hilus neurons (Spiegel et al., 2013). In early studies, global depletion of SST in rats by cysteamine hydrochloride, a somatostatin inhibiting agent, impaired learning and memory in a passive avoidance task (Yamazaki et al., 1996); and similar studies have shown impairment in spatial discrimination in mice (Guillou et al., 1998; Epelbaum et al., 2009; Tuboly and Vecsei, 2013). Interestingly, optogenetic inhibition of GABAergic DG hilus interneuron activity impairs spatial learning and memory retrieval (Andrews-Zwilling et al., 2012), although this effect was via the entire GABA interneuron population with no implicit selectivity for SST/GABA neurons. Multiple studies have reported that SST interneurons inhibit excitatory synaptic transmission excitability, resulting in silencing or hyperpolarization in hippocampus (see (Liguz-Lecznar et al., 2016) for review); and increases in inhibition (of hippocampal interneurons) is well-known to be an important cellular mechanism in learning and memory (McEchron and Disterhoft, 1997; Knott et al., 2002; Ruediger et al., 2011). In more recent studies, an enhancement in inhibition onto CA1 pyramidal neurons during learning in a trace eye blink conditioning task, was found to be mediated by an increase in intrinsic excitability of SST interneurons in mice and rats (McKay et al., 2013), and inactivating SST dendrite-targeting interneurons during aversive stimuli increased CA1 pyramidal cell population responses and prevented fear learning in mice (Lovett-Barron et al., 2014).

Therefore, our second goal was to assess the behavioral effects of Cre-recombinase-induced local deletion of RXFP3 in the DG hilus of adult floxed-RXFP3 mice. In these studies we observed a strong and consistent deficiency in the performance of AAV^(1/2)-Cre-IRES-eGFP-treated floxed-RXFP3 mice in an appetitive reference memory task in a T-maze. In a continuous spontaneous alternation Y-maze task, RXFP3-depleted mice made fewer alternations in the first min of the task, suggesting impairment in spatial working memory. In terms of the functional implications of these findings, it is proposed that the RLN3-containing inputs to the GABA/SST-positive neurons and other GABA neuron populations in the DG hilus can regulate hippocampal and cognitive processing. If this is the case, depletion of RXFP3 from GABA and/or GABA/SST neurons in the DG hilus may disrupt disinhibition of target neurons, and weaken synaptic inhibition of excitatory neurons, a key component in compromised cognitive processes. This, in turn, could produce impaired learning and memory, via effects within the directly targeted hippocampus and via remote, relayed effects in the MS/DB. Further studies are required to explore this hypothesis, including an examination of the effect of RXFP3 activation by validated RXFP3-specific agonist peptides (Liu et al., 2005; Shabanpoor et al., 2012; Zhang et al., 2015) on DG hilus SST neuron activity, perhaps in a convenient strain of SST reporter mice (Ma et al., 2006; Peng et al., 2013).

In contrast to these effects, however, a deficiency in spatial learning and long-term memory was not observed based on the performance of AAV^(1/2)-Cre-IRES-eGFP-treated floxed-RXFP3 mice in the MWM. The differences observed in these tests of memory retention suggest a specific role for hilar RLN3/RXFP3 signaling in learning and memory that may reflect the different neurochemical and psychological processes involved in acquiring memory. Optimal performance in the MWM requires the use of allocentric strategies where distal cues provide a geometric reference to an animal's current location and this form of memory consolidation is primarily hippocampal-dependent (Vorhees and Williams, 2006; Garthe et al., 2009). While effective performance in the appetitive T-maze task also requires the use of hippocampal-dependent allocentric strategies whereby the animal is rewarded for choosing a constant, designated goal arm on the basis of its spatial location (Deacon et al., 2002; Reisel et al., 2002; Sanderson et al., 2008; von Engelhardt et al., 2008), differences in memory consolidation between these two spatial reference memory tests have been highlighted (Hodges, 1996; Deacon et al., 2002; Bannerman et al., 2012). The most prominent refers to the distinct differences in the motivational aspect of both tasks. The MWM is motivated by the desire to escape onto a safe platform, whereas the appetitive T-maze is motivated by food reward. Therefore, differences in reinforcement may affect the strategy adopted and activate different brain circuits involved in motivation and reward. Furthermore, although both tests require the use of allocentric strategies, the appetitive T-maze requires the mouse to use spatial information to guide selection between two alternative responses and to learn to make a constant body turn for a food reward. Taken together, the results from the present study suggest RXFP3 in the DG hilus is not essential for spatial tasks that rely

solely on the use of allocentric spatial maps, but may influence tasks which integrate multiple strategies for using spatial information to guide selection between alternative responses that lead to a reinforcing reward. Moreover, in regard to the contradictory effects observed in the spontaneous alternation Y-maze task and MWM in the present study, spontaneous alternation has been shown to be a more reliable measure in detecting hippocampal dysfunction than the commonly used MWM (Rawlins and Olton, 1982; Reisel et al., 2002; Deacon and Rawlins, 2005). Although both tasks detect spatial memory deficits in mice with whole lesions of the hippocampus (Morris et al., 1982; Lalonde, 2002; Deacon and Rawlins, 2005), consistent with the results from the present study, a deficit in spatial memory following deletion of the GluR-A (GluR1) AMPA receptor subunit was only detected in a spontaneous alternation T-maze task, whereas spatial reference memory was not affected in a MWM (Reisel et al., 2002).

In a continuous SAT in a Y-maze, mice with RXFP3 depleted from the DG hilus displayed significantly lower rates of alternation (PAS, 36%) relative to control mice (PAS, 68%) in the first minute of the task, demonstrating an impairment in spatial working memory. After 3 min, DG hilus RXFP3 depleted mice still displayed low rates of alternation (PAS 47%), but the control mice had similar rates (PAS, 52%), suggesting poor spatial working memory performance in both treatment groups. However, the weak performance of the control group assessed over a 3-min session is not surprising. The continuous nature of the task means there is considerable inter-trial interference, a likely reason for low rates of performance in both T-maze (Gerlai et al., 1994) and Y-maze (Hsiao et al., 1996; Deacon et al., 2002) tasks (for review see, Deacon and Rawlins, 2006). Therefore, the decline in PAS observed over the subsequent 2 and 3 min time bins in both treatment groups is likely due to methodological issues associated with the task, whereby the first min is the period in which the most reliable measures are observed.

Lastly, RXFP3 depletion from the DG hilus did not alter anxiety-like behavior in an EPM or L/D box, tested 1 and 3 days, respectively, after the MWM test with its associated swim stress (Engelmann et al., 2006). In the mice studied, we observed AAV^(1/2) spread along the rostral-caudal length of the DG hilus and CA3, including part of the ventral hippocampus, which is functionally implicated in the control of emotion and affect, and wherein RLN3-positive nerve fibres and RXFP3 mRNA are abundant (Ma et al., 2009b; Smith et al., 2010). However, the viral injections in the present study did not cover the entire dorsoventral extent of the DG hilus in ventral hippocampus, reducing the likelihood of observing an effect on anxiety-like behavior. The RLN3/RXFP3 system in mice and rats has been shown to be involved in stress responses (Tanaka et al., 2005; Banerjee et al., 2010) and modulation of anxiety-like behavior (Ryan et al., 2013; Zhang et al., 2015), with pronounced changes in RLN3 mRNA expression and RLN3-IR levels in response to stressors (Tanaka et al., 2005; Lenglos et al., 2013; Ma et al., 2013; Walker et al., 2015; Calvez et al., 2016). Therefore, further studies with more extensive and/or focused depletion of RXFP3 in ventral hippocampus are warranted.

Conclusions

The anatomical distribution of the RLN3/RXFP3 system in the major nodes of the septohippocampal system (Ma et al., 2009c; Smith et al., 2010; Olucha-Bordonau et al., 2012) suggests it contributes to important neuronal circuits in cognition, and learning and memory. This first study to investigate the specific role of RLN3/RXFP3 signaling in the mouse hippocampus has demonstrated neuronal contacts with distinct populations of GABAergic neurons in the hippocampus, and that depletion of RXFP3 within the DG hilus can produce impairments in spatial reference and working memory **in an appetitive T-maze and spontaneous alternation task in a Y-maze, respectively**. These data suggest that the RLN3-containing pathway from the NI to the hippocampus can mediate inhibitory control of local and septal target neurons via RXFP3, and modulate memory processes.

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Figure Legends

FIGURE 1. RXFP3 mRNA in the mouse dorsal hippocampus.

Representative dark-field photomicrographs of emulsion autoradiographic analysis of RXFP3 mRNA expression in the hippocampus. **(A)** Silver grains for RXFP3 mRNA were detected in the DG hilus **(B)**, the pyramidal layer and stratum oriens of the CA1 **(C)** and the radiatum, pyramidal layer and stratum oriens of the CA3 **(D)**. Boxed areas in **(A)** indicate the areas illustrated in magnified images in **(B, C and D)**.

FIGURE 2. RLN3-positive nerve fibres and terminals in the mouse dorsal hippocampus.

Mosaic confocal image (20× magnification represented as z-stack maximum projections, 1 µm intervals) of mouse dorsal hippocampus illustrating RLN3-positive nerve fibres and terminals **(A)**. Nerve fibres and terminals containing RLN3-IR were detected in the DG hilus **(B)**, the pyramidal layer and stratum oriens of the CA1 **(C)** and the radiatum, pyramidal layer and stratum oriens on the CA3 **(D)**. Boxed areas in **(A)** indicate the areas illustrated in magnified images (63× maximum projections z-stacks, 0.1 µm intervals) **(B, C, D)**. Scale bars, 200 µm **(A)** and 20 µm **(B, C and D)**.

FIGURE 3. Comparative distribution in mouse dorsal hippocampus of RXFP3 mRNA and mRNAs encoding peptide and protein markers of hippocampal interneurons.

(A) Distribution of RXFP3 mRNA in adult mouse hippocampus with abundant expression detected in neurons located in CA1 stratum oriens, across the CA3 region and within the hilus of the dentate gyrus (DG) with higher magnification views of CA3 and DG (boxed areas) provided **(A', A'')**. The comparative distribution of GAD67 **(B)**, SST **(C)**, PV **(D)**, and CR **(E)** mRNAs illustrate the strong similarity between the distribution of RXFP3 mRNA positive neurons and *some* of the GAD67, SST, PV and CR mRNA positive neurons in the CA1 stratum oriens, CA3 region and hilus, suggesting that RXFP3 is expressed by a population of GABA interneurons (or projection neurons) in these areas. Images adapted from the Allen Brain Institute Gene Expression Atlas <www.brain-map.org>.

FIGURE 4. Comparative distribution of RLN3-immunoreactive nerve fibres and neurons positive for amino acid, peptide or protein markers of hippocampal interneurons in dorsal hippocampus of adult mouse.

Mosaic confocal images (20× magnification represented as z-stack maximum projections, 1 µm intervals) of mouse DG illustrating **(A)** RLN3 (red), SST (green) and PV (blue) immunoreactivity, **(B)** RLN3 (red) and GABA (green) immunoreactivity, and **(C)** RLN3 (red) and CR (green) immunoreactivity. Boxed areas in A-C are further illustrated (63× magnification z-stacks, 0.1 µm intervals) and represented as three-dimensional orthogonal maximum projections orientated in 3 planes (y/x, x/z and y/z) **(A', B', C')**. Dotted lines in these panels indicate appositions of RLN3-positive elements with hippocampal neurons in the y/x, x/z and y/z planes. Scale bars represent 100 µm **(A, B, C)** and 20 µm **(A', B', C')**.

FIGURE 5. Comparative distribution of eGFP, Cre-recombinase and somatostatin (SST) in hilar neurons of dorsal hippocampus in mice injected with the AAV^(1/2)-Cre-IRES-eGFP viral vector.

Bilateral injections of AAV^(1/2)-Cre-IRES-eGFP into the dentate gyrus (DG) of floxed-RXFP3 mice resulted in expression of eGFP **(A)** and Cre-immunoreactivity **(B)** in neurons of the DG hilus. Cre-immunoreactivity and endogenous eGFP immunofluorescence were consistently observed in SST-immunoreactive neurons in the region **(C, D)**, consistent with the likely deletion of any RXFP3 expression from these neurons. In the higher magnification images (inserts), arrowheads indicate some of many neurons in which eGFP, Cre and SST are co-localized. Scale bars, 100 μ m **(A-D)** and 20 μ m (inserts).

FIGURE 6. Effect of an AAV^(1/2)-Cre-IRES-eGFP or an AAV^(1/2)-eGFP viral vector injection into the dorsal dentate gyrus hilus of floxed-RXFP3 mice on levels of RXFP3 mRNA in the target and CA3 regions, and a schematic illustration of the virus distribution observed.

Representative dark-field photomicrographs of nuclear emulsion autoradiographic images of the distribution of RXFP3 mRNA in DG hilus of floxed RXFP3 mice labeled by an [³⁵S]-riboprobe (Lenglos et al., 2014) following bilateral injections of either **(A)** AAV^(1/2)-eGFP, or **(B)** AAV^(1/2)-Cre-IRES-eGFP. A significant reduction in relative RXFP3 mRNA levels was observed after bilateral AAV^(1/2)-Cre-IRES-eGFP injections into the DG hilus, relative to bilateral AAV^(1/2)-eGFP (control) injections in **(C)** hilus and **(D)** CA3 (unpaired t-test, ** $p < 0.01$, $n = 3 - 5$ mice per group). Data are expressed as mean $\pm$ SEM. **(E)** Schematic representation of viral spread observed within the DG of treated mice. The distribution of virus and Cre recombinase was similar on both sides of the hippocampus in all of the mice included in the final behavioural analysis. (Coronal brain images adapted from a stereotaxic mouse brain atlas (Paxinos and Franklin 2001)).

FIGURE 7. Effect of RXFP3 deletion in the dentate gyrus hilus of adult floxed-RXFP3 mice on subsequent spatial reference memory in an appetitive T-maze.

An AAV^(1/2)-Cre-IRES-eGFP or an AAV^(1/2)-eGFP (control) viral vector was injected bilaterally into the dentate gyrus (DG) of groups of floxed-RXFP3 mice and their spatial memory was tested after a 3-week recovery period. **(A)** During acquisition training, mice with RXFP3 deleted from DG hilus made significantly less correct choices than control mice on day 1-3 of testing in a hippocampus-dependent spatial reference memory task - an appetitive T-maze. RXFP3 deleted and control mice performed similarly during reversal training (2-way repeated measures ANOVA, Bonferroni post-hoc analysis, * $p < 0.05$). **(B)** Mice with RXFP3 deleted from DG hilus took significantly more days of training than control mice to meet criteria during acquisition (Mann-Whitney U-test, * $p < 0.05$). No treatment differences were observed during reversal training. Data are expressed as mean $\pm$ SEM, $n = 12 - 13$ mice per group.

FIGURE 8. Effect of RXFP3 deletion in the dentate gyrus hilus of adult floxed-RXFP3 mice on subsequent performance in a spontaneous alternation task in a Y-maze.

An AAV^(1/2)-Cre-IRES-eGFP or an AAV^(1/2)-eGFP (control) viral vector was injected bilaterally into the dentate gyrus (DG) of groups of floxed-RXFP3 mice and their performance in a spontaneous alternation task in a Y-maze was tested after a 3-week recovery period. Mice with RXFP3 deleted from DG hilus displayed a significant reduction in percent alternation score (PAS) in the first min of a spontaneous alternation task (SAT) in the Y-maze, relative to control mice. No treatment differences in PAS were observed during the 2nd or 3rd min bins of the test (Unpaired t-test, *** $p < 0.001$, $n = 12 - 13$ mice per group). Horizontal dotted line indicates 50% chance level.

FIGURE 9. Effect of RXFP3 deletion in the dentate gyrus hilus of adult floxed-RXFP3 mice on spatial learning and long-term reference memory in a Morris water maze (MWM) test.

An AAV^(1/2)-Cre-IRES-eGFP or an AAV^(1/2)-eGFP (control) viral vector was injected bilaterally into the dentate gyrus (DG) of groups of floxed-RXFP3 mice and their performance in a Morris water maze (MWM) test was tested after a 3-week recovery period. **(A)** Mice with RXFP3 deleted from DG hilus and matched control mice displayed normal spatial learning in the MWM. **(B)** On the probe day, both groups spent significantly more time in the target quadrant relative to other quadrants (one-way ANOVA, Bonferroni post-hoc analysis, * $p < 0.05$, **** $p < 0.0001$, $n = 12 - 13$ mice per group). Data are expressed as mean $\pm$ SEM. Horizontal dotted line indicate 25% chance level. Abbreviations: T, target quadrant (SW), A, adjacent quadrants (NW, SE) and O opposite quadrant (NE).

FIGURE 1. RXFP3 mRNA in the mouse dorsal hippocampus.

Representative dark-field photomicrographs of emulsion autoradiographic analysis of RXFP3 mRNA expression in the hippocampus. (A) Silver grains for RXFP3 mRNA were detected in the DG hilus (B), the pyramidal layer and stratum oriens of the CA1 (C) and the radiatum, pyramidal layer and stratum oriens of the CA3 (D). Boxed areas in (A) indicate the areas illustrated in magnified images in (B, C and D).

124x126mm (300 x 300 DPI)

FIGURE 2. RLN3-positive nerve fibres and terminals in the mouse dorsal hippocampus. Mosaic confocal image (20× magnification represented as z-stack maximum projections, 1 μ m intervals) of mouse dorsal hippocampus illustrating RLN3-positive nerve fibres and terminals (A). Nerve fibres and terminals containing RLN3-IR were detected in the DG hilus (B), the pyramidal layer and stratum oriens of the CA1 (C) and the radiatum, pyramidal layer and stratum oriens on the CA3 (D). Boxed areas in (A) indicate the areas illustrated in magnified images (63× maximum projections z-stacks, 0.1 μ m intervals) (B, C, D). Scale bars, 200 μ m (A) and 20 μ m (B, C and D).

151x187mm (300 x 300 DPI)

FIGURE 3. Comparative distribution in mouse dorsal hippocampus of RXFP3 mRNA and mRNAs encoding peptide and protein markers of hippocampal interneurons. (A) Distribution of RXFP3 mRNA in adult mouse hippocampus with abundant expression detected in neurons located in CA1 stratum oriens, across the CA3 region and within the hilus of the dentate gyrus (DG) with higher magnification views of CA3 and DG (boxed areas) provided (A', A''). The comparative distribution of GAD67 (B), SST (C), PV (D), and CR (E) mRNAs illustrate the strong similarity between the distribution of RXFP3 mRNA positive neurons and some of the GAD67, SST, PV and CR mRNA positive neurons in the CA1 stratum oriens, CA3 region and hilus, suggesting that RXFP3 is expressed by a population of GABA interneurons (or projection neurons) in these areas. Images adapted from the Allen Brain Institute Gene Expression Atlas <www.brain-map.org>.

109x91mm (300 x 300 DPI)

A

FIGURE 4. Comparative distribution of RLN3-immunoreactive nerve fibres and neurons positive for amino acid, peptide or protein markers of hippocampal interneurons in dorsal hippocampus of adult mouse. Mosaic confocal images (20× magnification represented as z-stack maximum projections, 1 μm intervals) of mouse DG illustrating (A) RLN3 (red), SST (green) and PV (blue) immunoreactivity, (B) RLN3 (red) and GABA (green) immunoreactivity, and (C) RLN3 (red) and CR (green) immunoreactivity. Boxed areas in A-C are further illustrated (63× magnification z-stacks, 0.1 μm intervals) and represented as three-dimensional orthogonal maximum projections orientated in 3 planes (y/x, x/z and y/z) (A', B', C'). Dotted lines in these panels indicate appositions of RLN3-positive elements with hippocampal neurons in the y/x, x/z and y/z planes. Scale bars represent 100 μm (A, B, C) and 20 μm (A', B', C').

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FIGURE 5. Comparative distribution of eGFP, Cre-recombinase and somatostatin (SST) in hilar neurons of dorsal hippocampus in mice injected with the AAV(1/2)-Cre-IRES-eGFP viral vector. Bilateral injections of AAV(1/2)-Cre-IRES-eGFP into the dentate gyrus (DG) of floxed-RXFP3 mice resulted in expression of eGFP (A) and Cre-immunoreactivity (B) in neurons of the DG hilus. Cre-immunoreactivity and endogenous eGFP immunofluorescence were consistently observed in SST-immunoreactive neurons in the region (C, D), consistent with the likely deletion of any RXFP3 expression from these neurons. In the higher magnification images (inserts), arrowheads indicate some of many neurons in which eGFP, Cre and SST are co-localized. Scale bars, 100 μ m (A-D) and 20 μ m (inserts).

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FIGURE 6. Effect of an AAV(1/2)-Cre-IRES-eGFP or an AAV(1/2)-eGFP viral vector injection into the dorsal dentate gyrus hilus of floxed-RXFP3 mice on levels of RXFP3 mRNA in the target and CA3 regions, and a schematic illustration of the virus distribution observed.

Representative dark-field photomicrographs of nuclear emulsion autoradiographic images of the distribution of RXFP3 mRNA in DG hilus of floxed RXFP3 mice labeled by an [³⁵S]-riboprobe (Lenglos et al., 2014) following bilateral injections of either (A) AAV(1/2)-eGFP, or (B) AAV(1/2)-Cre-IRES-eGFP. A significant reduction in relative RXFP3 mRNA levels was observed after bilateral AAV(1/2)-Cre-IRES-eGFP injections into the DG hilus, relative to bilateral AAV(1/2)-eGFP (control) injections in (C) hilus and (D) CA3 (unpaired t-test, ** $p < 0.01$, $n = 3 - 5$ mice per group). Data are expressed as mean $\pm$ SEM. (E) Schematic representation of viral spread observed within the DG of treated mice. The distribution of virus and Cre recombinase was similar on both sides of the hippocampus in all of the mice included in the final behavioural analysis. (Coronal brain images adapted from a stereotaxic mouse brain atlas (Paxinos and Franklin 2001)).

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A

FIGURE 7. Effect of RXFP3 deletion in the dentate gyrus hilus of adult floxed-RXFP3 mice on subsequent spatial reference memory in an appetitive T-maze.

An AAV(1/2)-Cre-IRES-eGFP or an AAV(1/2)-eGFP (control) viral vector was injected bilaterally into the dentate gyrus (DG) of groups of floxed-RXFP3 mice and their spatial memory was tested after a 3-week recovery period. (A) During acquisition training, mice with RXFP3 deleted from DG hilus made significantly less correct choices than control mice on day 1-3 of testing in a hippocampus-dependent spatial reference memory task - an appetitive T-maze. RXFP3 deleted and control mice performed similarly during reversal training (2-way repeated measures ANOVA, Bonferroni post-hoc analysis, * $p < 0.05$). (B) Mice with RXFP3 deleted from DG hilus took significantly more days of training than control mice to meet criteria during acquisition (Mann-Whitney U-test, * $p < 0.05$). No treatment differences were observed during reversal training. Data are expressed as mean $\pm$ SEM, $n = 12 - 13$ mice per group.

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FIGURE 8. Effect of RXFP3 deletion in the dentate gyrus hilus of adult floxed-RXFP3 mice on subsequent performance in a spontaneous alternation task in a Y-maze.

An AAV(1/2)-Cre-IRES-eGFP or an AAV(1/2)-eGFP (control) viral vector was injected bilaterally into the dentate gyrus (DG) of groups of floxed-RXFP3 mice and their performance in a spontaneous alternation task in a Y-maze was tested after a 3-week recovery period. Mice with RXFP3 deleted from DG hilus displayed a significant reduction in percent alternation score (PAS) in the first min of a spontaneous alternation task (SAT) in the Y-maze, relative to control mice. No treatment differences in PAS were observed during the 2nd or 3rd min bins of the test (Unpaired t-test, *** $p < 0.001$, $n = 12 - 13$ mice per group). Horizontal dotted line indicates 50% chance level.

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FIGURE 9. Effect of RXFP3 deletion in the dentate gyrus hilus of adult floxed-RXFP3 mice on spatial learning and long-term reference memory in a Morris water maze (MWM) test.

An AAV(1/2)-Cre-IRES-eGFP or an AAV(1/2)-eGFP (control) viral vector was injected bilaterally into the dentate gyrus (DG) of groups of floxed-RXFP3 mice and their performance in a Morris water maze (MWM) test was tested after a 3-week recovery period. (A) Mice with RXFP3 deleted from DG hilus and matched control mice displayed normal spatial learning in the MWM. (B) On the probe day, both groups spent significantly more time in the target quadrant relative to other quadrants (one-way ANOVA, Bonferroni post-hoc analysis, * $p < 0.05$, **** $p < 0.0001$, $n = 12 - 13$ mice per group). Data are expressed as mean $\pm$ SEM. Horizontal dotted line indicate 25% chance level. Abbreviations: T, target quadrant (SW), A, adjacent quadrants (NW, SE) and O opposite quadrant (NE).

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FIGURE S1. Comparative distribution of RLN3-immunoreactive nerve fibres and hippocampal neurons immunoreactive for somatostatin (SST) and parvalbumin (PV) in the dorsal DG hilus of adult mouse hippocampus.

Mosaic confocal images (20× magnification represented as z-stack maximum projections, 1 µm intervals) of mouse DG (reproduced from Figure 4), illustrating (A) RLN3 (red), SST (green) and PV (blue) immunoreactivity. Further details of the putative interactions between RLN3 inputs and SST neurons were recorded in this section. The DG hilar SST-positive neurons indicated by arrowheads in A, are illustrated at higher resolution in (B) and (C). There are several points of close apposition between the RLN3 fibres/boutons and the soma and proximal/distal processes of the SST neurons (yellow). Scale bars, 100 µm (A) and 20 µm (B, C).

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A

FIGURE S2. Comparative distribution of Cre-recombinase and somatostatin (SST) in DG hilar neurons of dorsal hippocampus in mice injected with the AAV(1/2)-Cre-IRES-eGFP viral vector. Bilateral injections of AAV(1/2)-Cre-IRES-eGFP into the DG of floxed-RXFP3 mice resulted in expression of Cre-immunoreactivity (red) (A, A') in SST neurons (blue) (B, B') of the DG hilus. Cre-immunoreactivity (pink) was consistently observed in the nuclei of SST-immunoreactive neurons in the region (C, C'), consistent with the deletion of RXFP3 expression from these neurons. In the higher magnification images of the boxed areas (A', B', C'), arrowheads indicate examples of neurons in which Cre and SST are co-localized. Scale bars, 100 μ m (A-C) and 20 μ m (A'-C').

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